

# Thermal and spectroscopic properties of light lanthanides (III) and sodium complexes of 2,5-pyridinedicarboxylic acid

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CCTA10 Special Issue

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**Abstract** Pyridine-2,5-dicarboxylic acid, known as isocinchomeric acid is one of six isomers containing two carboxylic groups. Light lanthanide (III) complexes with pyridine-2,5-dicarboxylic acid with general formula  $\text{Ln}_2\text{L}_3 \cdot n\text{H}_2\text{O}$ , where  $n = 8, 9$ , were obtained. Their thermal and spectroscopic properties were studied. Sodium salt was obtained as  $\text{Na}_2\text{L} \cdot \text{H}_2\text{O}$ . Hydrated complexes of La(III), Ce(III), Pr(III), Nd(III), Sm(III), Eu(III) and Gd(III) are stable to 313–333 K, whereas  $\text{Na}_2\text{L} \cdot \text{H}_2\text{O}$  is stable to about 333 K. Dehydration process for all compounds runs in one stage, next they decompose into appropriate lanthanide oxalates, oxocarbonates carbonates and finally to metal oxides. Bands of  $\nu\text{COOH}$  vibrations at 1736 and  $1728\text{ cm}^{-1}$  disappear on complex spectra and  $\nu_{\text{as}}$  and  $\nu_{\text{s}}$  of  $\text{COO}^-$  groups appear thus indicating that complexation process took place.

**Keywords** Lanthanide complexes ·  
Pyridinedicarboxylates · 2,5-Pyridinedicarboxylates ·  
Thermal decomposition

## Introduction

Rapid development of coordination polymers synthesis has been reported in recent years [1–3]. It is mainly caused by their potential applications and possibility of creating multidimensional structures required in biochemistry as well as in pharmacy. Due to the presence of various coordinating

atoms like O and N pyridinedicarboxylic acids are ligands which play an instant role in building such crystal architectures. Large number of transition and lanthanide complexes with series of the pyridinedicarboxylic acids (PDCA), which include the 2,3-, 2,4-, 2,5-, 2,6-, 3,4- and 3,5-pyridinedicarboxylic isomers, have been reported in the recent two decades. Although there are many publications about their complexes with transition metals, there are few papers about compounds formed by 2,5-pydcH<sub>2</sub> ligand. There is a wide range of coordination possibilities among isomers of isocinchomeric acid. They can create helicates and other supramolecular structures as well as nanostructures [4]. Isocinchomeric acid has beneficial structure to cage materials synthesis. It is mainly caused by the presence of two carboxylate groups separated by the nitrogen and carbon atom from the ring. Metal ion can be therefore easily coordinated and supramolecular and multidimensional structures can be created [5, 6]. Isocinchomeric acid can be herein promising ligand for MOFs structure building or cage materials where small molecules may be captured. Both the carboxylic groups can stabilize three-dimensional structures, and hence hydrogen bonds creation. It was reported that each carboxylic group can accept up to four hydrogen bonds [5, 6].

Hence due to the presence of five donor atoms in the molecule, pyridine-2,5-dicarboxylate anion has many different possibilities of coordination with lanthanide ions. Like Lewis base, it can also act as chelating or multidentate bridging ligand [7]. There were many compounds reported in which M–O and M–N coordination was observed [8–10]. Copper(II) ion in complex with 2,4-pyridinedicarboxylate anion was coordinated through nitrogen atom from the aromatic ring and one carboxylate anion situated next to the heteroatom, while the second carboxylic group became idle [11]. Similar coordination was observed for copper(II) complex with 3,4-pyridinedicarboxylic acid

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[12], zinc(II), nickel(II) [13], and cobalt(II) [14] complexes with 2,4-pyridinedicarboxylic acid. In mixed complexes, for example Ln(III)–Cu(II), transition metal ion is coordinated through nitrogen atom from the aromatic ring and carboxylate anion situated in second position, while the lanthanide ion is coordinated by the second carboxylate group (in fourth position) [15]. Therefore, it is highly possible that transition metal ions would be coordinated only by one carboxylate group and nitrogen atom, while lanthanide ions only through carboxylate anions in pyridinedicarboxylic acids. The presence of lone electron pair in the nitrogen atom from the aromatic ring and oxygen atoms from two different carboxylic groups cause the possibility of creating polymeric [16], dimeric [17] and rarely monomeric [18] structures. Lanthanides act as hard Pearson's acids; therefore, coordination through oxygen atoms from carboxylate groups mainly takes place. There are few examples where coordination between lanthanide cation and nitrogen atom from pyridine ring takes place. Geometric orientation of donor atoms in neodymium (III) complex with 2,6-pyridinedicarboxylic acid almost force lanthanide–nitrogen bonding [19]. It is mainly caused by the symmetrically spacing of carboxylic groups in 2,6-pyridinedicarboxylic acids. The chance of such lanthanide complexation among the other isomers of pyridinedicarboxylic acid, as well as isocinchomeric acid, is far lower, mainly due to steric conditions, although we can not exclude such possibility [20]. The lanthanide (III) coordination takes place through carboxylate groups, which are deprotonated in all complexes. Hence to the high density on oxygen atoms of these groups [21], there are many ways of metal ions coordination. Carboxylate group can have chelate [22], bridging, chelate-bridging [23] or monodentate. The geometry of these groups is often different what can be the reason of different ways of metal coordination [24]. Hence, many different spacial structures can be created. The best way to define the structure of organometallic compound is to obtain a monocrystal. We did not obtain the right crystal for analysis, though we concentrated on the IR spectra of obtained complexes analysis. IR spectra analyses were presented in this article and many possibilities of lanthanide ions coordination through donor atoms were discussed.

Although knowledge about biological meaning of pyridinedicarboxylates is still very poor, 2,5-pyridinedicarboxylic acid inhibits GA  $2\beta$ -hydroxylase and proline 4-hydroxylase enzymes [25]. It also accelerates the redox reactions between alcohols and chromic acid [26]. Pyridinedicarboxylic acids also act as chelating agents of chromium, zinc, manganese, copper, iron and molybdenum [7].

In this article, we report the synthesis, spectroscopic and thermal studies of obtained light lanthanide (III) complexes with isocinchomeric acid.

## Experimental

### Materials and methods

All chemicals employed were commercially available from Sigma and Aldrich Company and used without further purification. The purity of used acid was 98%. The IR spectra were recorded with 1725 $\times$  Perkin Elmer spectrometer at the range of 4000–200  $\text{cm}^{-1}$  using the KBr technique.

Thermogravimetric analyses were conducted on Setsys 16/18 analyzer in dynamic air atmosphere. Therefore, 6.96–7.51 mg samples were heated in the range of 303–1273 K in ceramic crucibles using a heating rate of 283 K  $\text{min}^{-1}$ . TG, DTG and DTA curves were registered.

Elementary analysis was made to determine composition and formulas of the obtained complexes. The analysis was carrying out using a CHN 2400 Perkin Elmer Analyzer. The water content was determined from the thermogravimetric curves of their mass loss and elementary analysis. The metal content was determined from TG curves.

Raman spectra were registered by Renishaw Raman Microscope via Reflex with 10 min exposure time and 785 nm laser edge.

Infrared and Raman theoretical calculations in DFTB3LYP/6-311++G\*\* were collected.

### Sample preparation

All compounds were obtained in double exchange method. To the hot solution of 0.1 M lanthanide chloride (pH ca. 5.3), stoichiometric amount of 0.1 M ammonium salt (pH ca. 5.8) of isocinchomeric acid was added dropwise, pH of final mixtures was about 5.0–5.3. Mixtures were heated on the electric stirrer for about 40 min at 338 K until full precipitation. Next, precipitates were filtered, washed with water to remove ammonium ions and dried at room temperature in desiccators to the stable mass. Cerium (III) complex was prepared from hexahydrated nitrate(V) with final pH 3.9.

Sodium salt was prepared from 1 M solution of sodium hydroxide into which 10% underflow of stoichiometric amount of 0.5 M 2,5-pyridinedicarboxylic acid solution was added dropwise. Final pH was equal to 7. The mixture was left on heater to remove excess of solvent. After 1 hour, it was left at room temperature for slower solvent evaporation.

Obtained complexes of lanthanum, europium and gadolinium were white, samarium was light cream, neodymium was light violet, praseodymium light green and cerium light yellow. Sodium salt was white.

## Results and discussion

### Elemental analysis

Carbon, hydrogen and nitrogen percentage contents determined for the obtained complexes are in agreement with found values (Table 1).

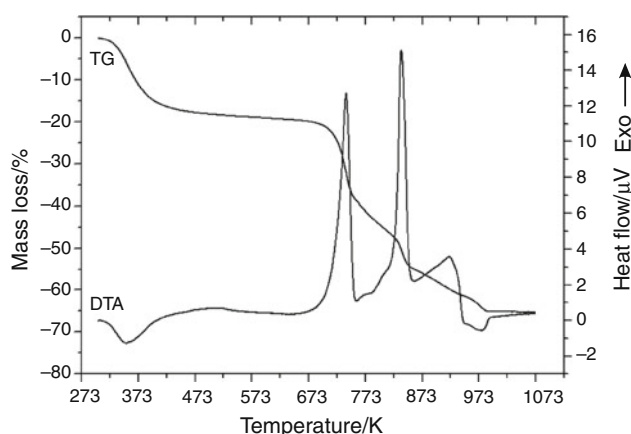
### Thermal analysis

Compounds obtained through the syntheses of light lanthanide ions with isocinchomeric acid were all hydrated with the general formula:  $\text{Ln}_2\text{L}_3 \cdot n\text{H}_2\text{O}$  where  $n$  is equal to 8 or 9, depending on the particular lanthanide complex.

The metal:ligand ratio in each complex is equal 2:3. Elementary analysis confirmed the ratio.

2,5-pyridinedicarboxylic acid is thermally stable up to 523 K when one stage decomposition process takes place (found mass loss: 99,091).

Monohydrated sodium salt was stable until 333 K when it decomposed into anhydrous complex  $\text{Na}_2\text{L}$ , which is stable up to 750 K. At 763 K, further decomposition takes place with sodium carbonate as the final decomposition product. Lanthanum complex was obtained as nonahydrated compound stable at 323 K (Fig. 1). All water molecules were lost in one stage dehydration. Anhydrous lanthanide (III) pyridinedicarboxylate is stable until 693 K, when it decomposes to oxalate, at 768 K to carbonate and at 873 to  $\text{La}_2\text{O}_3$ . Cerium (III) and praseodymium (III) complexes, both containing nine water molecules, decomposed similarly through anhydrous compounds to appropriate metal oxides  $\text{CeO}_2$  and  $\text{Pr}_6\text{O}_{11}$ . Both complexes started their dehydration at 313 K, although anhydrous praseodymium complex seem to be more stable with temperatures of decomposition at 713 K for Pr and 648 K for Ce. Octahydrated neodymium (III) complex, as well as the rest of obtained lanthanide complexes, was stable to 333 K. Dehydration process ran in one stage. Anhydrous compound was stable until 723 K and decomposed through carbonate, oxocarbonate to  $\text{Nd}_2\text{O}_3$ . Thermal decomposition



**Fig. 1** TG and DTA decomposition curves of  $\text{La}_2\text{L}_3 \cdot 9\text{H}_2\text{O}$

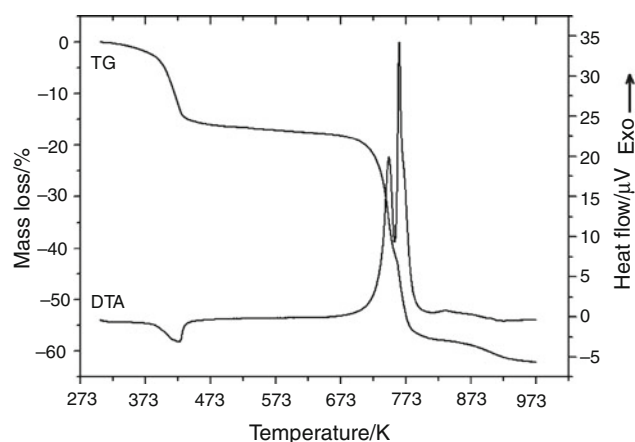
of  $\text{Sm}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$ ,  $\text{Eu}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$ ,  $\text{Gd}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$  (Fig. 2) ran similarly through three stages for each compound. First dehydration process which began at 313 K for Sm and at 323 K for Eu and Gd. Anhydrous complexes are stable until 718 K (Sm), 673 K (Eu) and 723 K (Gd) when decomposition to oxocarbonates takes place. Calculated and found mass losses were presented in Table 2.

### IR and Raman analysis

Hence to the difficulties with ascribing particular frequencies to bands on infrared spectra, theoretical IR and Raman calculations for acid were registered. These difficulties were mostly connected with bands corresponding to the ring vibrations region, which contains stretching vibrations of C=N bands from pyridine ring. According to the reported literature data, this band occurs at about 1290 and 1120  $\text{cm}^{-1}$  [27] and at 3490–3450  $\text{cm}^{-1}$  [28]. Hence to the theoretical data, we can assume that there are no single specific frequencies for C=N vibrations, because they are always connected with stretching or bending vibrations of pyridine ring. Stretching vibrations of C–H and C=N bands occur at 1637 and 1608  $\text{cm}^{-1}$  on theoretical IR spectrum. At the range of highest frequencies,

**Table 1** Results of elementary analysis of 2,5-pyridinedicarboxylates of light lanthanide complexes

Complexes	C%		H%		N%	
	Calculated	Found	Calculated	Found	Calculated	Found
$\text{La}_2\text{L}_3 \cdot 9\text{H}_2\text{O}$	26.95	26.54	2.88	3.07	4.49	4.36
$\text{Ce}_2\text{L}_3 \cdot 9\text{H}_2\text{O}$	26.87	26.35	2.88	3.02	4.48	4.30
$\text{Pr}_2\text{L}_3 \cdot 9\text{H}_2\text{O}$	26.84	26.28	2.87	3.14	4.47	4.32
$\text{Nd}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$	27.17	26.38	2.69	2.86	4.53	4.31
$\text{Sm}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$	26.81	26.42	2.66	2.87	4.47	4.42
$\text{Eu}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$	26.72	26.14	2.65	2.84	4.45	4.44
$\text{Gd}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$	26.43	25.86	2.62	2.81	4.40	4.40



**Fig. 2** TG and DTA decomposition curves of  $\text{Gd}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$

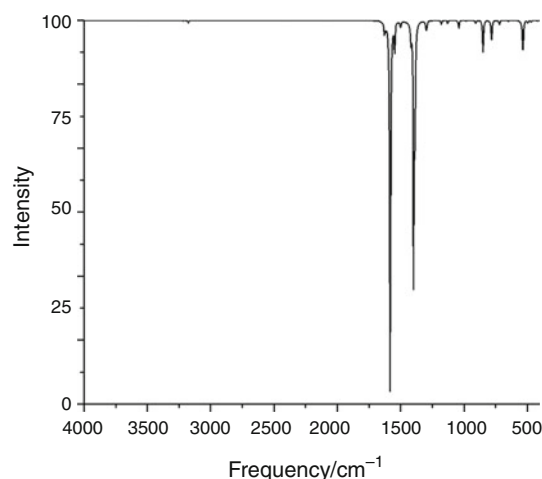
**Table 2** Thermal data of light lanthanides (III) complexes and sodium salt with 2,5-pyridinedicarboxylic acid

Compound	Range of decomposition temp./K	Mass loss/%		Stable products of decomposition
		Calculated	Found	
$\text{Na}_2\text{L}_3 \cdot \text{H}_2\text{O}$	333–423	7.86	8.07	$\text{Na}_2\text{L}_3$
	763–848	53.71	52.35	$\text{Na}_2\text{CO}_3$
$\text{La}_2\text{L}_3 \cdot 9\text{H}_2\text{O}$	323–453	17.32	17.59	$\text{La}_2\text{L}_3$
	693–768	42.06	41.50	$\text{La}_2(\text{O}_x)_3$
	768–873	51.05	51.89	$\text{La}_2(\text{CO}_3)_3$
	873–923	65.16	65.04	$\text{La}_2\text{O}_3$
$\text{Ce}_2\text{L}_3 \cdot 9\text{H}_2\text{O}$	313–433	17.28	16.80	$\text{Ce}_2\text{L}_3$
	648–718	63.31	63.55	$\text{CeO}_2$
$\text{Pr}_2\text{L}_3 \cdot 9\text{H}_2\text{O}$	313–443	17.25	17.01	$\text{Pr}_2\text{L}_3$
	713–833	63.75	63.00	$\text{Pr}_6\text{O}_{11}$
$\text{Nd}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$	333–453	15.52	15.66	$\text{Nd}_2\text{L}_3$
	723–773	49.51	48.83	$\text{Nd}_2(\text{CO}_3)_3$
	773–873	59.00	60.03	$\text{Nd}_2\text{O}_2\text{CO}_3$
	873–953	63.73	63.54	$\text{Nd}_2\text{O}_3$
$\text{Sm}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$	313–468	15.32	15.11	$\text{Sm}_2\text{L}_3$
	718–793	58.23	58.08	$\text{Sm}_2\text{O}_2\text{CO}_3$
	793–948	62.90	62.99	$\text{Sm}_2\text{O}_3$
$\text{Eu}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$	323–463	15.27	15.40	$\text{Eu}_2\text{L}_3$
	673–788	58.03	58.60	$\text{Eu}_2\text{O}_2\text{CO}_3$
	788–943	62.69	62.80	$\text{Eu}_2\text{O}_3$
$\text{Gd}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$	323–453	15.10	15.22	$\text{Gd}_2\text{L}_3$
	723–798	57.38	56.85	$\text{Gd}_2\text{O}_2\text{CO}_3$
	798–948	61.99	61.57	$\text{Gd}_2\text{O}_3$

stretching vibrations of OH groups from carboxylic group of fifth and second position, respectively, are present. Frequencies at 3198, 3199 and 3216  $\text{cm}^{-1}$  correspond to the stretching vibrations of hydrogen atoms C–H from aromatic ring. Bands at 1828 and 1797  $\text{cm}^{-1}$  correspond to the  $\nu_{\text{as}}\text{COO}$  vibrations in second and fifth position,

respectively. We can therefore assume that higher frequency value for COO group in second position is caused by some interaction between carboxylic groups placed nearest nitrogen atom from aromatic ring. At lower frequencies, different ring vibrations, breathing and bending, are present. Experimental IR spectrum of 2,5-pyridinedicarboxylic acid was interpreted on the basis of theoretical spectrum. At the range of highest frequencies, weak bands of hydroxyl group stretching vibrations  $\nu\text{OH}$  are present as well as quite strong bands of hydrogen vibrations from aromatic ring. At about 1736  $\text{cm}^{-1}$ , strong splitted band of carboxylic groups is present. At the range of 1624–1536  $\text{cm}^{-1}$ , there are three medium bands of aromatic ring stretching vibrations, which also appear at 1408 and 1384  $\text{cm}^{-1}$ . At the range of lower frequencies (1328–360  $\text{cm}^{-1}$ ), numerous bands of ring vibrations are present with the most characteristic at 1328  $\text{cm}^{-1}$  ( $\beta$  C–H)<sub>ar</sub>, 1012  $\text{cm}^{-1}$  ( $\gamma$  C–H in 3 and 4 position), 932  $\text{cm}^{-1}$  ( $\gamma$  C–H in 6 position) and 896  $\text{cm}^{-1}$  corresponding to the breathing ring vibrations. On theoretical IR spectrum of sodium salt (Fig. 3), two bands of high intensity can be observed at about 1400 and 1580  $\text{cm}^{-1}$ . These bands correspond to symmetric and asymmetric vibrations of deprotonated carboxylic groups, respectively. At the range of lower frequencies, there were observed five more bands corresponding to the different ring vibrations. Frequencies for acid spectrum and all obtained light lanthanide (III) complexes spectra were collected in Table 3.

Experimental IR spectra of sodium salt (Fig. 4) and light lanthanide complexes (Figs. 5, 6) with isocinchomeric acid are very similar. At the range of highest frequencies, bands of stretching  $\nu\text{OH}$  group from water molecules, vibrations are present. On each complexes spectra, we can observe strong bands at about 1600  $\text{cm}^{-1}$ , which correspond to the stretching asymmetric vibrations of carboxylate groups. Bands of  $\nu_{\text{sym}}\text{COO}^-$  vibrations occur at 1408, 1364  $\text{cm}^{-1}$  (sodium salt), 1396, 1364  $\text{cm}^{-1}$  (La), 1400, 1364  $\text{cm}^{-1}$



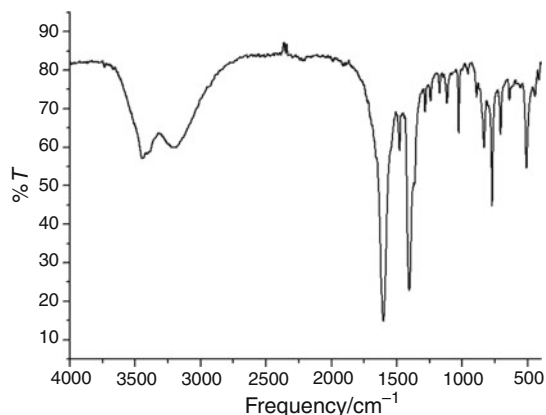
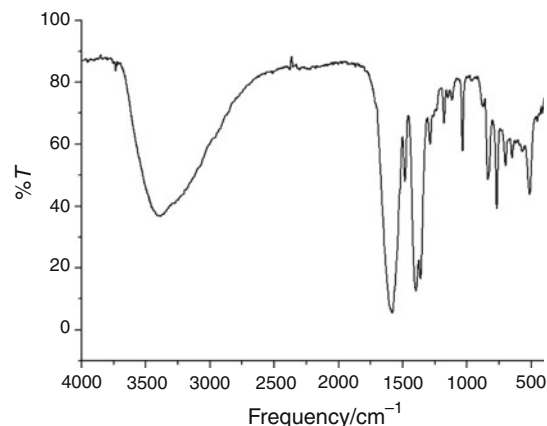
**Fig. 3** Theoretical IR spectrum of sodium salt

**Table 3** Frequencies of maximum of absorption bands in IR and Raman (experimental) spectra of 2,5-pyridinedicarboxylic acid and pyridinedicarboxylates of light lanthanides and Na/cm<sup>-1</sup>

Complex	$\nu\text{OH}$		$\nu\text{COOH}$		$\nu_{\text{as}}\text{COO}^-$		$\nu_{\text{s}}\text{COO}^-$		Ring vibrations $\nu$ , $\beta$ , $\gamma$ + C=N		$\nu\text{Me-O}$	
	IR	Raman	IR	Raman	IR	Raman	IR	Raman	IR	Raman	IR	Raman
C <sub>7</sub> H <sub>5</sub> NO <sub>4</sub>	3440	3680, 3648	1736, 1728	–	–	1797, 1828	–	–	1624, 1596, 1328	1608, 1628	–	–
Na <sub>2</sub> L·1H <sub>2</sub> O	3448	–	–	–	1600	–	1408, 1364	–	1284–640	1285–641	508	–
La <sub>2</sub> L <sub>3</sub> ·9H <sub>2</sub> O	3392	–	–	–	1584	–	1396, 1364	–	1288–648	1597–651	512	–
Ce <sub>2</sub> L <sub>3</sub> ·9H <sub>2</sub> O	3400	–	–	–	1580	–	1400, 1364	–	1288–572	1597–651	512	–
Pr <sub>2</sub> L <sub>3</sub> ·9H <sub>2</sub> O	3400	–	–	–	1584	–	1400, 1364	–	1288–658	1597–652	516	–
Nd <sub>2</sub> L <sub>3</sub> ·8H <sub>2</sub> O	3388	–	–	–	1584	–	1400, 1364	–	1288–1120	1643–1205	524	–
Sm <sub>2</sub> L <sub>3</sub> ·8H <sub>2</sub> O	3380	–	–	–	1588	–	1408, 1364	–	1288–652	1596–650	524	–
Eu <sub>2</sub> L <sub>3</sub> ·8H <sub>2</sub> O	3416	–	–	–	1588	–	1400, 1364	–	1288–624	1596–650	524	–
Gd <sub>2</sub> L <sub>3</sub> ·8H <sub>2</sub> O	3368	–	–	–	1604, 1556	–	1400, 1364	–	1288–652	1596–650	524	–

L = C<sub>5</sub>H<sub>3</sub>N(COO<sup>-</sup>)<sub>2</sub>

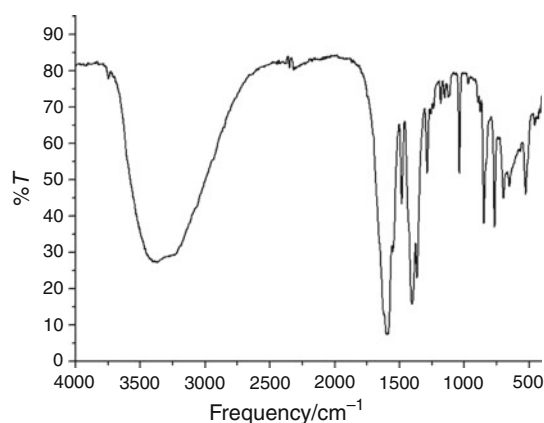
(Ce, Pr), 1400, 1364 cm<sup>-1</sup> (Nd, Eu, Gd), 1408, 1364 cm<sup>-1</sup> (Sm). Only bands of europium (III) and gadolinium (III) complexes are little splitted. There is no bands characteristic for carboxylic group from the acid at about 1700 cm<sup>-1</sup>, what indicates that coordination processes took place in each compound. We can therefore assume that different ways of metal coordination may take place. At the range of 1288–640 cm<sup>-1</sup>, various bands of stretching, bending, wagging and out of plane ring vibrations are present on all complexes spectra. Medium intensity bands characteristic for breathing ring vibrations occur at 832 cm<sup>-1</sup> (Na), 840 cm<sup>-1</sup> (La, Pr), 836 cm<sup>-1</sup> (Ce), 848 cm<sup>-1</sup> (Nd, Sm, Eu) and 852 cm<sup>-1</sup> (Gd). Also, medium intensity bands

**Fig. 4** Experimental IR spectrum of sodium salt**Fig. 5** IR spectrum of La<sub>2</sub>L<sub>3</sub>·9H<sub>2</sub>O

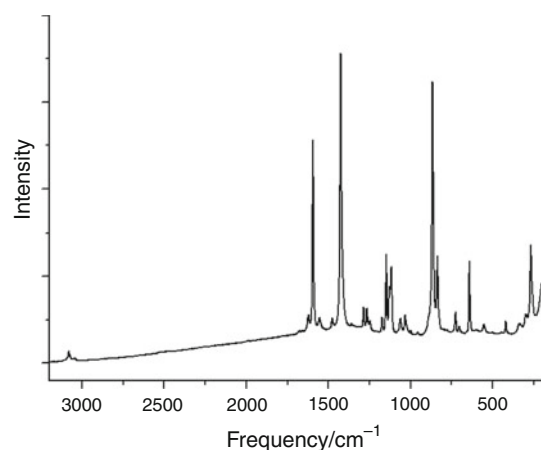
corresponding to metal–ligand coordination appear at the range of lowest frequencies, i.e., 508 cm<sup>-1</sup> (Na), 512 cm<sup>-1</sup> (La, Ce), 516 cm<sup>-1</sup> (Pr) and 524 cm<sup>-1</sup> (Nd, Sm, Eu, Gd).

Raman Spectra were registered for all obtained complexes. Experimental spectra of sodium, lanthanum and gadolinium complexes were presented in Figs. 7, 8 and 9, respectively. The most intensive bands are present at about 1597, 880, and 1440 cm<sup>-1</sup> which correspond to stretching ring vibrations ( $\nu\text{C-H}$ ,  $\nu\text{C=N}$ ,  $\nu\text{C-C}$ ) and bending  $\beta$  C–O vibrations, respectively. They are much more intensive than bands present on IR spectra. Breathing ring vibrations with medium intensity bands occur at 830 cm<sup>-1</sup>.

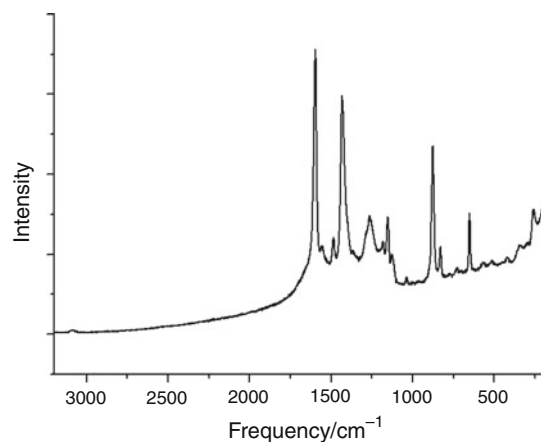




**Fig. 6** IR spectrum of  $\text{Gd}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$



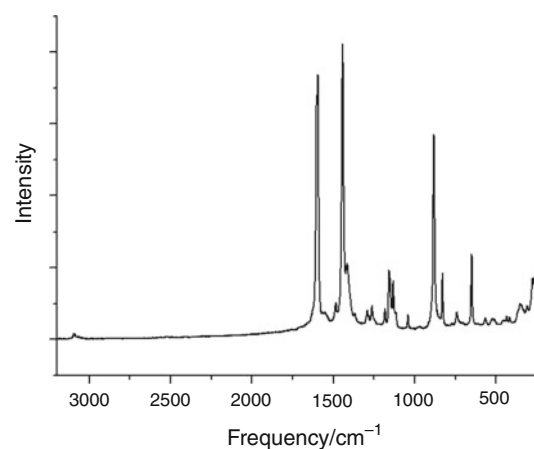
**Fig. 7** Experimental Raman spectrum of sodium salt



**Fig. 8** Raman spectrum of  $\text{La}_2\text{L}_3 \cdot 9\text{H}_2\text{O}$

## Conclusions

Light lanthanide (III) and sodium complexes with 2,5-pyridinedicarboxylic acid have been synthesized by the



**Fig. 9** Raman spectrum of  $\text{Gd}_2\text{L}_3 \cdot 8\text{H}_2\text{O}$

double exchange method. Based on IR and Raman analysis, possible ways of coordination have been investigated and discussed. Coordination probably takes place through carboxylate groups from 2,5-pyridinedicarboxylic ring, although we cannot exclude the possibility of coordination through nitrogen atom. Theoretical IR calculations made for isocinchomeric acid indicate on the interaction between nitrogen atom and carboxylic group placed in the second position.

Light lanthanides complexes with isocinchomeric acid are very good potential materials for large area microporous compounds synthesis. These hydrated compounds are thermally stable up to 313–333 K, and during heated process, they lose all water molecules in one stage with creation of compounds with strictly defined composition. Anhydrous complexes are thermally stable up to 673–763 K. Thermal stability of compound structure is necessary condition for their potential application as microporous compounds.

This stage of our work allowed us to precisely identify the stability and spectroscopic properties of the obtained complexes. Examination of these complexes will be continuing in the direction of obtaining microporous materials and studies on their sorption properties. Our further work will be connected with MOF-like structure synthesis in hydrothermal conditions.

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